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(54) **Arginyl tRNA synthase**

(57) The invention provides argS polypeptides and DNA (RNA) encoding argS polypeptides and methods

for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing argS polypeptides to screen for antibacterial compounds.

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**Description****RELATED APPLICATIONS**

This application claims benefit of GB application number 9619071.5, filed September 12, 1996.

**FIELD OF THE INVENTION**

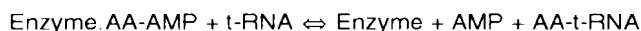
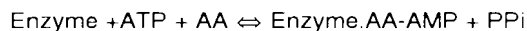
This invention relates to newly identified polynucleotides and polypeptides, and their production and uses, as well as their variants, agonists and antagonists, and their uses. In particular, in these and in other regards, the invention relates to novel polynucleotides and polypeptides of the arginyl tRNA synthetase family, hereinafter referred to as "argS".

**BACKGROUND OF THE INVENTION**

The Streptococci make up a medically important genera of microbes known to cause several types of disease in humans, including, for example, otitis media, conjunctivitis, pneumonia, bacteremia, meningitis, sinusitis, pleural empyema and endocarditis, and most particularly meningitis, such as for example infection of cerebrospinal fluid. Since its isolation more than 100 years ago, *Streptococcus pneumoniae* has been one of the more intensively studied microbes. For example, much of our early understanding that DNA is, in fact, the genetic material was predicated on the work of Griffith and of Avery, Macleod and McCarty using this microbe. Despite the vast amount of research with *S. pneumoniae*, many questions concerning the virulence of this microbe remain. It is particularly preferred to employ Streptococcal genes and gene products as targets for the development of antibiotics.

The frequency of *Streptococcus pneumoniae* infections has risen dramatically in the past 20 years. This has been attributed to the emergence of multiply antibiotic resistant strains and an increasing population of people with weakened immune systems. It is no longer uncommon to isolate *Streptococcus pneumoniae* strains which are resistant to some or all of the standard antibiotics. This has created a demand for both new anti-microbial agents and diagnostic tests for this organism.

t-RNA synthetases have a primary role in protein synthesis according to the following scheme:



in which AA is an amino acid.

Inhibition of this process leads to a reduction in the levels of charged t-RNA and this triggers a cascade of responses known as the stringent response, the result of which is the induction of a state of dormancy in the organism. As such selective inhibitors of bacterial t-RNA synthetase have potential as antibacterial agents. One example of such is mupirocin which is a selective inhibitor of isoleucyl t-RNA synthetase. Other t-RNA synthetases are now being examined as possible anti-bacterial targets, this process being greatly assisted by the isolation of the synthetase.

Clearly, there is a need for factors, such as the novel compounds of the invention, that have a present benefit of being useful to screen compounds for antibiotic activity. Such factors are also useful to determine their role in pathogenesis of infection, dysfunction and disease. There is also a need for identification and characterization of such factors and their antagonists and agonists which can play a role in preventing, ameliorating or correcting infections, dysfunctions or diseases.

The polypeptides of the invention have amino acid sequence homology to a known *Saccharomyces cerevisiae* mitochondrial arginyl tRNA synthetase protein.

**SUMMARY OF THE INVENTION**

It is an object of the invention to provide polypeptides that have been identified as novel argS polypeptides by homology between the amino acid sequence set out in Table 1 [SEQ ID NO: 2] and a known amino acid sequence or sequences of other proteins such as *Saccharomyces cerevisiae* mitochondrial arginyl tRNA synthetase protein.

It is a further object of the invention to provide polynucleotides that encode argS polypeptides, particularly polynucleotides that encode the polypeptide herein designated argS.

In a particularly preferred embodiment of the invention the polynucleotide comprises a region encoding argS

polypeptides comprising the sequence set out in Table 1 [SEQ ID NO:1] which includes a full length gene, or a variant thereof.

In another particularly preferred embodiment of the invention there is a novel argS protein from *Streptococcus pneumoniae* comprising the amino acid sequence of Table 1 [SEQ ID NO:2], or a variant thereof.

In accordance with another aspect of the invention there is provided an isolated nucleic acid molecule encoding a mature polypeptide expressible by the *Streptococcus pneumoniae* 0100993 strain contained in the deposited strain.

A further aspect of the invention there are provided isolated nucleic acid molecules encoding argS, particularly *Streptococcus pneumoniae* argS, including mRNAs, cDNAs, genomic DNAs. Further embodiments of the invention include biologically, diagnostically, prophylactically, clinically or therapeutically useful variants thereof, and compositions comprising the same.

In accordance with another aspect of the invention, there is provided the use of a polynucleotide of the invention for therapeutic or prophylactic purposes, in particular genetic immunization. Among the particularly preferred embodiments of the invention are naturally occurring allelic variants of argS and polypeptides encoded thereby.

Another aspect of the invention there are provided novel polypeptides of *Streptococcus pneumoniae* referred to herein as argS as well as biologically, diagnostically, prophylactically, clinically or therapeutically useful variants thereof, and compositions comprising the same.

Among the particularly preferred embodiments of the invention are variants of argS polypeptide encoded by naturally occurring alleles of the argS gene.

In a preferred embodiment of the invention there are provided methods for producing the aforementioned argS polypeptides.

In accordance with yet another aspect of the invention, there are provided inhibitors to such polypeptides, useful as antibacterial agents, including, for example, antibodies.

In accordance with certain preferred embodiments of the invention, there are provided products, compositions and methods for assessing argS expression, treating disease, for example, otitis media, conjunctivitis, pneumonia, bacteremia, meningitis, sinusitis, pleural empyema and endocarditis, and most particularly meningitis, such as for example infection of cerebrospinal fluid, assaying genetic variation, and administering a argS polypeptide or polynucleotide to an organism to raise an immunological response against a bacteria, especially a *Streptococcus pneumoniae* bacteria.

In accordance with certain preferred embodiments of this and other aspects of the invention there are provided polynucleotides that hybridize to argS polynucleotide sequences, particularly under stringent conditions.

In certain preferred embodiments of the invention there are provided antibodies against argS polypeptides.

In other embodiments of the invention there are provided methods for identifying compounds which bind to or otherwise interact with and inhibit or activate an activity of a polypeptide or polynucleotide of the invention comprising: contacting a polypeptide or polynucleotide of the invention with a compound to be screened under conditions to permit binding to or other interaction between the compound and the polypeptide or polynucleotide to assess the binding to or other interaction with the compound, such binding or interaction being associated with a second component capable of providing a detectable signal in response to the binding or interaction of the polypeptide or polynucleotide with the compound; and determining whether the compound binds to or otherwise interacts with and activates or inhibits an activity of the polypeptide or polynucleotide by detecting the presence or absence of a signal generated from the binding or interaction of the compound with the polypeptide or polynucleotide.

In accordance with yet another aspect of the invention, there are provided argS agonists and antagonists, preferably bacteriostatic or bacteriocidal agonists and antagonists.

In a further aspect of the invention there are provided compositions comprising a argS polynucleotide or a argS polypeptide for administration to a cell or to a multicellular organism.

Various changes and modifications within the spirit and scope of the disclosed invention will become readily apparent to those skilled in the art from reading the following descriptions and from reading the other parts of the present disclosure.

## GLOSSARY

The following definitions are provided to facilitate understanding of certain terms used frequently herein.

"Host cell" is a cell which has been transformed or transfected, or is capable of transformation or transfection by an exogenous polynucleotide sequence.

"Identity," as known in the art, is a relationship between two or more polypeptide sequences or two or more polynucleotide sequences, as determined by comparing the sequences. In the art, "identity" also means the degree of sequence relatedness between polypeptide or polynucleotide sequences, as the case may be, as determined by the match between strings of such sequences. "Identity" and "similarity" can be readily calculated by known methods, including but not limited to those described in (*Computational Molecular Biology*, Lesk, A.M., ed., Oxford University Press, New York, 1988; *Biocomputing: Informatics and Genome Projects*, Smith, D.W., ed., Academic Press, New

York, 1993; *Computer Analysis of Sequence Data*, Part I, Griffin, A.M., and Griffin, H.G., eds., Humana Press, New Jersey, 1994; *Sequence Analysis in Molecular Biology*, von Heinje, G., Academic Press, 1987; and *Sequence Analysis Primer*, Gribskov, M. and Devereux, J., eds., M Stockton Press, New York, 1991; and Carillo, H., and Lipman, D., *SIAM J. Applied Math.*, 48: 1073 (1988). Preferred methods to determine identity are designed to give the largest match between the sequences tested. Methods to determine identity and similarity are codified in publicly available computer programs. Preferred computer program methods to determine identity and similarity between two sequences include, but are not limited to, the GCG program package (Devereux, J., et al., *Nucleic Acids Research* 12(1): 387 (1984)), BLASTP, BLASTN, and FASTA (Atschul, S.F. et al., *J. Molec. Biol.* 215: 403-410 (1990)). The BLAST X program is publicly available from NCBI and other sources (*BLAST Manual*, Altschul, S., et al., NCBI NLM NIH Bethesda, MD 20894; Altschul, S., et al., *J. Mol. Biol.* 215: 403-410 (1990)). As an illustration, by a polynucleotide having a nucleotide sequence having at least, for example, 95% "identity" to a reference nucleotide sequence of SEQ ID NO: 1 it is intended that the nucleotide sequence of the polynucleotide is identical to the reference sequence except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence of SEQ ID NO: 1. In other words, to obtain a polynucleotide having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted or substituted with another nucleotide, or a number of nucleotides up to 5% of the total nucleotides in the reference sequence may be inserted into the reference sequence. These mutations of the reference sequence may occur at the 5 or 3 terminal positions of the reference nucleotide sequence or anywhere between those terminal positions, interspersed either individually among nucleotides in the reference sequence or in one or more contiguous groups within the reference sequence. Analogously, by a polypeptide having an amino acid sequence having at least, for example, 95% identity to a reference amino acid sequence of SEQ ID NO: 2 is intended that the amino acid sequence of the polypeptide is identical to the reference sequence except that the polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the reference amino acid of SEQ ID NO: 2. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a reference amino acid sequence, up to 5% of the amino acid residues in the reference sequence may be deleted or substituted with another amino acid, or a number of amino acids up to 5% of the total amino acid residues in the reference sequence may be inserted into the reference sequence. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

"Isolated" means altered "by the hand of man" from its natural state, i.e., if it occurs in nature, it has been changed or removed from its original environment, or both. For example, a polynucleotide or a polypeptide naturally present in a living organism is not "isolated," but the same polynucleotide or polypeptide separated from the coexisting materials of its natural state is "isolated", as the term is employed herein.

"Polynucleotide(s)" generally refers to any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. "Polynucleotide(s)" include, without limitation, single- and double-stranded DNA, DNA that is a mixture of single- and double-stranded regions or single-, double- and triple-stranded regions, single- and double-stranded RNA, and RNA that is mixture of single- and double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded, or triple-stranded regions, or a mixture of single- and double-stranded regions. In addition, "polynucleotide" as used herein refers to triple-stranded regions comprising RNA or DNA or both RNA and DNA. The strands in such regions may be from the same molecule or from different molecules. The regions may include all of one or more of the molecules, but more typically involve only a region of some of the molecules. One of the molecules of a triple-helical region often is an oligonucleotide. As used herein, the term "polynucleotide(s)" also includes DNAs or RNAs as described above that contain one or more modified bases. Thus, DNAs or RNAs with backbones modified for stability or for other reasons are "polynucleotide(s)" as that term is intended herein. Moreover, DNAs or RNAs comprising unusual bases, such as inosine, or modified bases, such as tritylated bases, to name just two examples, are polynucleotides as the term is used herein. It will be appreciated that a great variety of modifications have been made to DNA and RNA that serve many useful purposes known to those of skill in the art. The term "polynucleotide(s)" as it is employed herein embraces such chemically, enzymatically or metabolically modified forms of polynucleotides, as well as the chemical forms of DNA and RNA characteristic of viruses and cells, including, for example, simple and complex cells. "Polynucleotide(s)" also embraces short polynucleotides often referred to as oligonucleotide(s).

"Polypeptide(s)" refers to any peptide or protein comprising two or more amino acids joined to each other by peptide bonds or modified peptide bonds. "Polypeptide(s)" refers to both short chains, commonly referred to as peptides, oligopeptides and oligomers and to longer chains generally referred to as proteins. Polypeptides may contain amino acids other than the 20 gene encoded amino acids. "Polypeptide(s)" include those modified either by natural processes, such as processing and other post-translational modifications, but also by chemical modification techniques. Such modifications are well described in basic texts and in more detailed monographs, as well as in a voluminous research literature, and they are well known to those of skill in the art. It will be appreciated that the same type of modification

may be present in the same or varying degree at several sites in a given polypeptide. Also, a given polypeptide may contain many types of modifications. Modifications can occur anywhere in a polypeptide, including the peptide backbone, the amino acid side-chains, and the amino or carboxyl termini. Modifications include, for example, acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphatidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine, formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, proteolytic processing, phosphorylation, prenylation, racemization, glycosylation, lipid attachment, sulfation, gamma-carboxylation of glutamic acid residues, hydroxylation and ADP-ribosylation, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins, such as arginylation, and ubiquitination. See, for instance, *PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES*, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993) and Wold, F., *Posttranslational Protein Modifications: Perspectives and Prospects*, pgs. 1-12 in *POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS*, B. C. Johnson, Ed., Academic Press, New York (1983); Seifter et al., *Meth. Enzymol.* 182:626-646 (1990) and Rattan et al., *Protein Synthesis: Posttranslational Modifications and Aging*, Ann. N.Y. Acad. Sci. 663: 48-62 (1992). Polypeptides may be branched or cyclic, with or without branching. Cyclic, branched and branched circular polypeptides may result from post-translational natural processes and may be made by entirely synthetic methods, as well.

"Variant(s)" as the term is used herein, is a polynucleotide or polypeptide that differs from a reference polynucleotide or polypeptide respectively, but retains essential properties. A typical variant of a polynucleotide differs in nucleotide sequence from another, reference polynucleotide. Changes in the nucleotide sequence of the variant may or may not alter the amino acid sequence of a polypeptide encoded by the reference polynucleotide. Nucleotide changes may result in amino acid substitutions, additions, deletions, fusions and truncations in the polypeptide encoded by the reference sequence, as discussed below. A typical variant of a polypeptide differs in amino acid sequence from another, reference polypeptide. Generally, differences are limited so that the sequences of the reference polypeptide and the variant are closely similar overall and, in many regions, identical. A variant and reference polypeptide may differ in amino acid sequence by one or more substitutions, additions, deletions in any combination. A substituted or inserted amino acid residue may or may not be one encoded by the genetic code. A variant of a polynucleotide or polypeptide may be a naturally occurring such as an allelic variant, or it may be a variant that is not known to occur naturally. Non-naturally occurring variants of polynucleotides and polypeptides may be made by mutagenesis techniques, by direct synthesis, and by other recombinant methods known to skilled artisans.

## DESCRIPTION OF THE INVENTION

The invention relates to novel argS polypeptides and polynucleotides as described in greater detail below. In particular, the invention relates to polypeptides and polynucleotides of a novel argS of *Streptococcus pneumoniae*, which is related by amino acid sequence homology to *Saccharomyces cerevisiae* mitochondrial arginyl tRNA synthetase polypeptide. The invention relates especially to argS having the nucleotide and amino acid sequences set out in Table 1 [SEQ ID NO: 1] and Table 1 [SEQ ID NO: 2] respectively, and to the argS nucleotide sequences of the DNA in the deposited strain and amino acid sequences encoded thereby.

TABLE 1

## argS Polynucleotide and Polypeptide Sequences

(A) Sequences from *Streptococcus pneumoniae* argS polynucleotide sequence [SEQ ID NO:1].

5'-1 ATGAATACAA AAGGATTGAT TGCTAGCGAA TTGGTTAGCA TCATTGATAG  
51 CATGGACCAA GAGGTAATTT TAAAGTTACT GGAAACCCCT AAAAATCAG  
101 AAATGGGGGA CATCGCTTTC CCTGCTTTTT CTCTTGCCAA AGTCGAACGT  
151 AAAGCACCAC AAATGATTGC GGCTAAACTG GCTGAAAAAA TGAACAGCCA  
201 AGCCTTTGAA AAAGTTGTCG CAACAGGACC TTACGTTAAC TTTTTCCTTG  
251 ATAAATCTGC CATTTCTGCT CAAGTATTGC AAGCTGTTAC CACTGAAAAA  
301 GAACACTATG CTGACCAAAA TATTGGTAAA CAAGAAAATG TTGTTATCGA  
351 CATGTCTAGT CCGAATATCG CTAAACCATT TTTTATTGGC CACCTGCGTT  
401 CAACTGTTAT CGGAGATAGC TTGTCACATA TTTTCCAAAA AATCGGTTAT  
451 CAAACGGTCA AGGTCAACCA TTTGGGAGAC TGGGGTAAAC AATTTGGGAT  
501 GTTGATTGTT GCCTACAAAA AATGGGGCGA CGAAGAAGCT GTAAAAGCTC  
551 ATCCAATCGA TGAATCCTT AAATCTATG TCCGCATCAA CGCTGAAGCT

601 GAAAATGACC CTAGCTTGA TTANGAAGCG CGCGAATGGT TCCGTAAACT  
 5 651 TGAAAATGGA GATGAGGAAG CTCTCGCTCT TTGGCAATGG TTCCGCGATG  
 701 AAAGTTTAGT GGAATTTAAC CGCCTTTACA ATGAATTGAA GGTGAATTT  
 10 751 GACAGCTATA ACGGAGAAGC CTTCTACAAT GATAAGATGG ATGCAGTTGT  
 801 AGACATTCTT TCTGAAAAAG GACTACTTCT TGAATCAGAA GGTGCCCAAG  
 15 851 TTGTCAATCT TGAGAAATAC GGAATTGAAC ATCCAGCTCT CATCAAGAAA  
 901 TCTGATGGTG CAACTCTCTA TATCACACGT GACTTGGCTG CAGCCCTTTA  
 20 951 CCGTAAAAAC GAATACGAAT TTGCTAAATC TATCTATGTC GTTGGTCAAG  
 1001 AACAATCTGC CCACTTTAAA CAGCTCAAAG CTGTCTTGCA AGAGATGGGC  
 25 1051 TACGACTGGA GTGACGACAT TACTCACGTT CCTTTTGGTT TGGTTACAAA  
 1101 AGAAGGGAAG AAACCTCTCTA CTCGTAAAGG GAATGTCATC TTGCTAGAGC  
 30 1151 CTACTGTTGC AGAGGCTGTT AGCCGTGCCA AGGTCCAAAT CGAGGCTAAA  
 35 1201 AATCCTGAAC TAGAAAACAA AGACCAAGTA GCACATGCTG TTGGGGTTGG  
 1251 AGCCATTAAA TTCTATGACC TCAAAACCGA CCGTACAAAT GGATACGACT  
 40 1301 TCGACCTAGA GGCTATGGTA TCCTTCGAGG GTGAAACTGG ACCTTACGTT  
 1351 CAATATGCCT ACGCTCGTAT CCAATCTATC TTACGCAAAG CCGATTTCAA  
 45 1401 ACCAGAAACA GCTGGCAACT ATAGCTTGAA TGATACTGAA AGCTGGGAAA  
 1451 TCATTAAACT CATTCAAGAC TTCCCACGTA TTATCAACCG TGCGGCAGAT  
 50 1501 AACTTTGAAC CTTCTATCAT TGCTAAATTT GCAATTAGCC TAGCTCAATC  
 55 1551 CTTTAACAAA TACTATGCAC ATACACGTAT CTTGGATGAA AGCCCAGAAC

1601 GCGACAGCCG TCTAGCCCTC AGCTACGCAA CCGCAGTCGT TCTCAAAGAA

5 1651 GCCCTTCGCT TGCTTGGAGT AGAAGCGCCA GAGAAGATGT AA-3'

(B) argS polypeptide sequence deduced from the polynucleotide sequence in this table  
[SEQ ID NO:2].

NH<sub>2</sub>-1 MNTKGLIASE LVSIIDSMDQ EVILKLETP KNSEMGDIAF PAFSLAKVER

51 KAPQMIAAKL AEKMNSQAFE KVVATGPYVN FFLDKSAISA QVLQAVTTEK

101 EHYADQNIGK QENVVIDMSS PNIAKPFFIG HLRSTVIGDS LSHIFQKIGY

151 QTVKVNHLGD WGKQFGMLIV AYKKWGDEEA VKAHPIDELL KLYVRINAEA

201 ENDPSLDXEA REWFRKLENG DEEALALWQW FRDESLVEFN RLYNELKVEF

251 DSYNGEAFYN DKMDAVVDIL SEKGLLLESE GAQVVNLEKY GIEHPALIKK

301 SDGATLYITR DLAAALYRKN EYEFAKSIYV VGQEQSAHFK QLKAVLQEMG

351 YDWSDDITHV PFGLVTKEGK KLSTRKGNVI LLEPTVAEAV SRAKVQIEAK

401 NPELENKDQV AHAVGVGAIK FYDLKTDRTN GYDFDLEAMV SFEGETGPYV

451 QYAYARIQSI LRKADFKPET AGNYSLNDTE SWEIIKLIQD FPRIINRAAD

501 NFEPSIIAKF AISLAQSFNK YYAHTRILDE SPERDSRLAL SYATAVVLKE

551 ALRLLGVEAP EKM-COOH

(C) Polynucleotide sequence embodiments [SEQ ID NO:1].

X-(R<sub>1</sub>)<sub>n-1</sub> ATGAATACAA AAGGATTGAT TGCTAGCGAA TTGGTTAGCA TCATTGATAG

51 CATGGACCAA GAGGTAATTT TAAAGTTACT GGAAACCCCT AAAAACTCAG

101 AAATGGGGGA CATCGCTTTC CCTGCTTTT CTCTTGCCAA AGTCGAACGT

151 AAAGCACCAC AAATGATTGC GGCTAAACTG GCTGAAAAAA TGAACAGCCA



201 AGCCTTTGAA AAAGTTGTCG CAACAGGACC TTACGTTAAC TTTTTCCTTG  
5 251 ATAAATCTGC CATTTCTGCT CAAGTATTGC AAGCTGTTAC CACTGAAAAA  
301 GAACACTATG CTGACCAAAA TATTGGTAAA CAAGAAAATG TTGTTATCGA  
10 351 CATGTCTAGT CCGAATATCG CTAAACCATT TTTTATTGGC CACCTGCGTT  
401 CAACTGTTAT CGGAGATAGC TTGTCACATA TTTTCCAAAA AATCGGTTAT  
15 451 CAAACGGTCA AGGTCAACCA TTTGGGAGAC TGGGGTAAAC AATTTGGGAT  
501 GTTGATTGTT GCCTACAAAA AATGGGGCGA CGAAGAAGCT GTAAAAGCTC  
20 551 ATCCAATCGA TGAACCTCTT AAACCTCTATG TCCGCATCAA CGCTGAAGCT  
25 601 GAAAATGACC CTAGCTTGGA TTANGAAGCG CGCGAATGGT TCCGTAAACT  
651 TGAAAATGGA GATGAGGAAG CTCTCGCTCT TTGGCAATGG TTCCGCGATG  
30 701 AAAGTTTAGT GGAATTTAAC CGCCTTTACA ATGAATTGAA GGTTGAATTT  
751 GACAGCTATA ACGGAGAAGC CTTCTACAAT GATAAGATGG ATGCAGTTGT  
35 801 AGACATTCTT TCTGAAAAAG GACTACTTCT TGAATCAGAA GGTGCCCCAAG  
851 TTGTCAATCT TGAGAAATAC GGAATTGAAC ATCCAGCTCT CATCAAGAAA  
40 901 TCTGATGGTG CAACTCTCTA TATCACACGT GACTTGGCTG CAGCCCTTTA  
951 CCGTAAAAAC GAATACGAAT TTGCTAAATC TATCTATGTC GTTGGTCAAG  
45 1001 AACAACTCTGC CCACTTTAAA CAGCTCAAAG CTGTCTTGCA AGAGATGGGC  
1051 TACGACTGGA GTGACGACAT TACTCACGTT CCTTTTGGTT TGTTACAAA  
50 1101 AGAAGGGAAG AAACCTCTCTA CTCGTAAAGG GAATGTCATC TTGCTAGAGC  
55 1151 CTA CTGTTGC AGAGGCTGTT AGCCGTGCCA AGGTCCAAAT CGAGGCTAAA

1201 AATCCTGAAC TAGAAAACAA AGACCAAGTA GCACATGCTG TTGGGGTTGG

5 1251 AGCCATTAAA TTCTATGACC TCAAAACCGA CCGTACAAAT GGATACGACT

1301 TCGACCTAGA GGCTATGGTA TCCTTCGAGG GTGAAACTGG ACCTTACGTT

10 1351 CAATATGCCT ACGCTCGTAT CCAATCTATC TTACGCAAAG CCGATTTCOA

1401 ACCAGAAACA GCTGGCAACT ATAGCTTGAA TGATACTGAA AGCTGGGAAA

15 1451 TCATTAAACT CATTCAAGAC TTCCCACGTA TTATCAACCG TCGGCGAGAT

1501 AACTTTGAAC CTTCTATCAT TGCTAAATTT GCAATTAGCC TAGCTCAATC

20 1551 CTTTAACAAA TACTATGCAC ATACACGTAT CTTGGATGAA AGCCCAGAAC

1601 GCGACAGCCG TCTAGCCCTC AGCTACGCAA CCGCAGTCGT TCTCAAAGAA

25 1651 GCCCTTCGCT TGCTTGGAGT AGAAGCGCCA GAGAAGATGT AA-(R<sub>2</sub>)<sub>n</sub>-Y

(D) Polypeptide sequence embodiments [SEQ ID NO:2].

30 X-(R<sub>1</sub>)<sub>n</sub>-1 MNTKGLIASE LVSIIDSMDQ EVILKLLETP KNSEMGDIAF PAFSLAKVER

51 KAPQMIAAKL AEKMNSQAFE KVVATGPYVN FFLDKSAISA QVLQAVTTEK

35 101 EHYADQNIGK QENVVIDMSS PNIAPFFIG HLRSTVIGDS LSHIFQKIGY

151 QTVKVNHLGD WGKQFGMLIV AYKKWGDEEA VKAHPIDELL KLYVRINAEA

40 201 ENDPSLDXEA REWFRKLENG DEEALALWQW FRDESLVEFN RLYNELKVEF

45 251 DSYNGEAFYN DKMDAVVDIL SEKGLLESE GAQVVNLEKY GIEHPALIKK

301 SDGATLYITR DLAAALYRKN EYEFAKSIYV VGQEQSAHFK QLKAVLQEMG

50 351 YDWSDDITHV PFGLVTKEGK KLSTRKGNVI LLEPTVAEAV SRAKVQIEAK

401 NPELENKDQV AHAVGVGAIK FYDLKTDRTN GYDFDLEAMV SFEGETGPYV

55 451 QYAYARIQSI LRKADFKPET AGNYSLNDE SWEIKLIQD FPRIINRAAD

501 NFEPSIIAKF AISLAQSFNK YYAHTRILDE SPERDSRLAL SYATAVVLKE

5 551 ALRLLGVEAP EKM- $(R_2)_n$ -Y

(E) Sequences from *Streptococcus pneumoniae* argS polynucleotide ORF sequence  
[SEQ ID NO:3].

5' -1 TTACGTTAAC TTTTCCTTG ATAAATCTGC CATTTCTGCT CAAGTATTGC

51 AAGCTGTTAC CACTGAAAAA GAACACTATG CTGACCAAAA TATTGGTAAA

101 CAAGAAAATG TTGTTATCGA CATGTCTAGT CCGAATATCG CTAAACCATT

151 TTTTATTGGC CACCTGCGTT CAACTGTTAT CGGAGATAGC TTGTCACATA

201 TTTTCCAAAA AATCGGTTAT CAAACGGTCA AGGTCAACCA TTTGGGAGAC

251 TGGGGTAAAC AATTTGGGAT GTTGATTGTT GCCTACAAAA AATGGGGCGA

301 CGAAGAAGCT GTAAAAGCTC ATCCAATCGA TGAATCCTT AAACCTCTATG

351 TCCGCATCAA CGCTGAAGCT GAAAATGACC CTAGCTTGGA TTANGAAGCG

401 CGCGAATGGT TCCGTAAACT TGAAAATGGA GATGAGGAAG CTCTCGCTCT

451 TTGGCAATGG TTCCGCGATG AAAGTTTAGT GGAATTTAAC CGCCTTTACA

501 ATGAATTGAA GGTTGAATTT GACAGCTATA ACGGAGAAGC CTTCTACAAT

551 GATAAGATGG ATGCAGTTGT AGACATTCTT TCTGAAAAAG GACTACTTCT

601 TGAATCAGAA GGTGCCCAAG TTGTCAATCT TGAGAAATAC GGAATTGAAC

651 ATCCAGCTCT CATCAAGAAA TCTGATGGTG CAACTCTCTA TATCACACGT

701 GACTTGGCTG CAGCCCTTTA CCGTAAAAAC GAATACGAAT TTGCTAAATC

751 TATCTATGTC GTTGGTCAAG AACAACTCTGC CCACTTTAAA CAGCTCAAAG

801 CTGTCTTGCA AGAGATGGGC TACGACTGGA GTGACGACAT TACTCACGTT

851 CCTTTTGGTT TGGTTACAAA AGAAGGGAAG AACTCTCTA CTCGTAAAGG  
 5 901 GAATGTCATC TTGCTAGAGC CTA CTGTTGC AGAGGCTGTT AGCCGTGCCA  
 951 AGGTCCAAAT CGAGGCTAAA AATCCTGAAC TAGAAAACAA AGACCAAGTA  
 10 1001 GCACATGCTG TTGGGGTTGG AGCCATTAAA TTCTATGACC TCAAACCGA  
 1051 CCGTACAAAT GGATACGACT TCGACCTAGA GGCTATGGTA TCCTTCGAGG  
 15 1101 GTGAAACTGG ACCTTACGTT CAATATGCCT ACGCTCGTAT CCAATCTATC  
 1151 TTACGCAAAG CCGATTTCOA ACCAGAAACA GCTGGCAACT ATAGCTTGAA  
 20 1201 TGATACTGAA AGCTGGGAAA TCATTAAACT CATTCAAGAC TTCCCACGTA  
 1251 TTATCAACCG TCGGCAGAT AACTTTGAAC CTTCTATCAT TGCTAAATTT  
 25 1301 GCAATTAGCC TAGCTCAATC CTTTAACAAA TACTATGCAC ATACACGTAT  
 1351 CTTGGATGAA AGCCCAGAAC GCGACAGCCG TCTAGCCCTC AGCTACGCAA  
 30 1401 CCGCAGTCGT TCTCAAAGAA GCCCTTCGCT TGCTTGAGT AGAAGCGCCA  
 35 1451 GAGAAGATGT AA-3'

(F) argS polypeptide sequence deduced from the polynucleotide ORF sequence in this table [SEQ ID NO:4].

40 NH<sub>2</sub>-1 YVNFFLDKSA ISAQVLQAVT TEKEHYADQN IGKQENVVID MSSPNIKPF  
 51 FIGHLRSTVI GDSLSHIFQK IGYQTVKVNH LGDWGKQFGM LIVAYKKWGD  
 45 101 EEAVKAHPID ELLKLYVRIN AEAENDPSLD XEAREWFRKL ENGDEEALAL  
 151 WQWFRDESLV EFNRLYNELK VEFDSYNGEA FYNDKMDAVV DILSEKGLLL  
 50 201 ESEGAQVVNL EKYGIEHPAL IKKSDGATLY ITRDLAAALY RKNEYEFAKS  
 55 251 IYVVGQEQSA HFKQLKAVLQ EMGYDWSDDI THVPFGLVTK EGKKLSTRKG

301 NVILLEPTVA EAVSRKVQI EAKNPELENK DQVAHAVGVG AIKFYDLKTD

5 351 RTNGYDFDLE AMVSFEGETG PYVQYAYARI QSILRKADFK PETAGNYSLN

401 DTESWEI IKL IQDFPRIINR AADNFEPSII AKFAISLAQS FNKYAHTRI

10 451 LDESPERDSR LALSATAVV LKEALRLLGV EAPEKM-COOH

## 15 Deposited materials

A deposit containing a *Streptococcus pneumoniae* 0100993 strain has been deposited with the National Collections of Industrial and Marine Bacteria Ltd. (herein "NCIMB"), 23 St. Machar Drive, Aberdeen AB2 1RY, Scotland on 11 April 1996 and assigned deposit number 40794. The deposit was described as *Streptococcus pneumoniae* 0100993 on deposit. On 17 April 1996 a *Streptococcus pneumoniae* 0100993 DNA library in *E. coli* was similarly deposited with the NCIMB and assigned deposit number 40800. The *Streptococcus pneumoniae* strain deposit is referred to herein as "the deposited strain" or as "the DNA of the deposited strain."

The deposited strain contains the full length *argS* gene. The sequence of the polynucleotides contained in the deposited strain, as well as the amino acid sequence of the polypeptide encoded thereby, are controlling in the event of any conflict with any description of sequences herein.

The deposit of the deposited strain has been made under the terms of the Budapest Treaty on the International Recognition of the Deposit of Micro-organisms for Purposes of Patent Procedure. The strain will be irrevocably and without restriction or condition released to the public upon the issuance of a patent. The deposited strain is provided merely as convenience to those of skill in the art and is not an admission that a deposit is required for enablement, such as that required under 35 U.S.C. §112.

A license may be required to make, use or sell the deposited strain, and compounds derived therefrom, and no such license is hereby granted.

## 35 Polypeptides

The polypeptides of the invention include the polypeptide of Table I [SEQ ID NO:2] (in particular the mature polypeptide) as well as polypeptides and fragments, particularly those which have the biological activity of *argS*, and also those which have at least 70% identity to a polypeptide of Table 1 [SEQ ID NOS:2 and 4] or the relevant portion, preferably at least 80% identity to a polypeptide of Table 1 [SEQ ID NOS:2 and 4], and more preferably at least 90% similarity (more preferably at least 90% identity) to a polypeptide of Table I [SEQ ID NOS:2 and 4] and still more preferably at least 95% similarity (still more preferably at least 95% identity) to a polypeptide of Table 1 [SEQ ID NOS:2 and 4] and also include portions of such polypeptides with such portion of the polypeptide generally containing at least 30 amino acids and more preferably at least 50 amino acids.

The invention also includes polypeptides of the formula set forth in Table 1 (D) [SEQ ID NO:2] wherein, at the amino terminus, X is hydrogen, and at the carboxyl terminus, Y is hydrogen or a metal, R<sub>1</sub> and R<sub>2</sub> is any amino acid residue, and n is an integer between 1 and 1000. Any stretch of amino acid residues denoted by either R group, where R is greater than 1, may be either a heteropolymer or a homopolymer, preferably a heteropolymer.

A fragment is a variant polypeptide having an amino acid sequence that entirely is the same as part but not all of the amino acid sequence of the aforementioned polypeptides. As with *argS* polypeptides fragments may be "fine-standing," or comprised within a larger polypeptide of which they form a part or region, most preferably as a single continuous region, a single larger polypeptide.

Preferred fragments include, for example, truncation polypeptides having a portion of an amino acid sequence of Table 1 [SEQ ID NOS:2 and 4], or of variants thereof, such as a continuous series of residues that includes the amino terminus, or a continuous series of residues that includes the carboxyl terminus. Degradation forms of the polypeptides of the invention in a host cell, particularly a *Streptococcus pneumoniae*, are also preferred. Further preferred are fragments characterized by structural or functional attributes such as fragments that comprise alpha-helix and alpha-helix forming regions, beta-sheet and beta-sheet-forming regions, turn and turn-forming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions,

surface-forming regions, substrate binding region, and high antigenic index regions.

Also preferred are biologically active fragments which are those fragments that mediate activities of argS, including those with a similar activity or an improved activity, or with a decreased undesirable activity. Also included are those fragments that are antigenic or immunogenic in an animal, especially in a human. Particularly preferred are fragments comprising receptors or domains of enzymes that confer a function essential for viability of *Streptococcus pneumoniae* or the ability to initiate, or maintain cause disease in an individual, particularly a human.

Variants that are fragments of the polypeptides of the invention may be employed for producing the corresponding full-length polypeptide by peptide synthesis; therefore, these variants may be employed as intermediates for producing the full-length polypeptides of the invention.

### Polynucleotides

Another aspect of the invention relates to isolated polynucleotides, including the full length gene, that encode the argS polypeptide having a deduced amino acid sequence of Table 1 [SEQ ID NOS:2 and 4] and polynucleotides closely related thereto and variants thereof.

Using the information provided herein, such as a polynucleotide sequence set out in Table 1 [SEQ ID NOS:1 and 3], a polynucleotide of the invention encoding argS polypeptide may be obtained using standard cloning and screening methods, such as those for cloning and sequencing chromosomal DNA fragments from bacteria using *Streptococcus pneumoniae* 0100993 cells as starting material, followed by obtaining a full length clone. For example, to obtain a polynucleotide sequence of the invention, such as a sequence given in Table 1 [SEQ ID NOS: 1 and 3], typically a library of clones of chromosomal DNA of *Streptococcus pneumoniae* 0100993 in *E. coli* or some other suitable host is probed with a radiolabeled oligonucleotide, preferably a 17-mer or longer, derived from a partial sequence. Clones carrying DNA identical to that of the probe can then be distinguished using stringent conditions. By sequencing the individual clones thus identified with sequencing primers designed from the original sequence it is then possible to extend the sequence in both directions to determine the full gene sequence. Conveniently, such sequencing is performed using denatured double stranded DNA prepared from a plasmid clone. Suitable techniques are described by Maniatis, T., Fritsch, E.F. and Sambrook et al., *MOLECULAR CLONING, A LABORATORY MANUAL*, 2nd Ed.; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (1989). (see in particular Screening By Hybridization 1.90 and Sequencing Denatured Double-Stranded DNA Templates 13.70). Illustrative of the invention, the polynucleotide set out in Table 1 [SEQ ID NO: 1] was discovered in a DNA library derived from *Streptococcus pneumoniae* 0100993.

The DNA sequence set out in Table I [SEQ ID NOS:1] contains an open reading frame encoding a protein having about the number of amino acid residues set forth in Table 1 [SEQ ID NO:2] with a deduced molecular weight that can be calculated using amino acid residue molecular weight values well known in the art. The polynucleotide of SEQ ID NO: 1, between nucleotide number 1 through number 1689 encodes the polypeptide of SEQ ID NO:2. The stop codon begins at nucleotide number 1690 of SEQ ID NO:1.

argS of the invention is structurally related to other proteins of the arginyl tRNA synthetase family, as shown by the results of sequencing the DNA encoding argS of the deposited strain. The protein exhibits greatest homology to *Saccharomyces cerevisiae* mitochondrial arginyl tRNA synthetase protein among known proteins. argS of Table I [SEQ ID NO:2] has about 35% identity over its entire length and about 62% similarity over its entire length with the amino acid sequence of *Saccharomyces cerevisiae* mitochondrial arginyl tRNA synthetase polypeptide.

The invention provides a polynucleotide sequence identical over its entire length to the coding sequence in Table 1 [SEQ ID NO:1]. Also provided by the invention is the coding sequence for the mature polypeptide or a fragment thereof, by itself as well as the coding sequence for the mature polypeptide or a fragment in reading frame with other coding sequence, such as those encoding a leader or secretory sequence, a pre-, or pro- or prepro- protein sequence. The polynucleotide may also contain non-coding sequences, including for example, but not limited to non-coding 5' and 3' sequences, such as the transcribed, non-translated sequences, termination signals, ribosome binding sites, sequences that stabilize mRNA, introns, polyadenylation signals, and additional coding sequence which encode additional amino acids. For example, a marker sequence that facilitates purification of the fused polypeptide can be encoded. In certain embodiments of the invention, the marker sequence is a hexa-histidine peptide, as provided in the pQE vector (Qiagen, Inc.) and described in Gentz *et al.*, *Proc. Natl. Acad. Sci., USA* 86: 821-824 (1989), or an HA tag (Wilson *et al.*, *Cell* 37: 767 (1984). Polynucleotides of the invention also include, but are not limited to, polynucleotides comprising a structural gene and its naturally associated sequences that control gene expression.

A preferred embodiment of the invention is a polynucleotide of comprising nucleotide 1 to 1689 or 1690 set forth in SEQ ID NO: 1 of Table 1 which encode the argS polypeptide.

The invention also includes polynucleotides of the formula set forth in Table 1 (C)[SEQ ID NO: 1] wherein, at the 5' end of the molecule, X is hydrogen, and at the 3' end of the molecule, Y is hydrogen or a metal, R<sub>1</sub> and R<sub>2</sub> is any nucleic acid residue, and n is an integer between 1 and 1000. Any stretch of nucleic acid residues denoted by either

R group, where R is greater than 1, may be either a heteropolymer or a homopolymer, preferably a heteropolymer.

The term "polynucleotide encoding a polypeptide" as used herein encompasses polynucleotides that include a sequence encoding a polypeptide of the invention, particularly a bacterial polypeptide and more particularly a polypeptide of the *Streptococcus pneumoniae* argS having the amino acid sequence set out in Table 1 [SEQ ID NO:2]. The term also encompasses polynucleotides that include a single continuous region or discontinuous regions encoding the polypeptide (for example, interrupted by integrated phage or an insertion sequence or editing) together with additional regions, that also may contain coding and/or non-coding sequences.

The invention further relates to variants of the polynucleotides described herein that encode for variants of the polypeptide having the deduced amino acid sequence of Table 1 [SEQ ID NO:2]. Variants that are fragments of the polynucleotides of the invention may be used to synthesize full-length polynucleotides of the invention.

Further particularly preferred embodiments are polynucleotides encoding argS variants, that have the amino acid sequence of argS polypeptide of Table 1 [SEQ ID NO:2] in which several, a few, 5 to 10, 1 to 5, 1 to 3, 2, 1 or no amino acid residues are substituted, deleted or added, in any combination. Especially preferred among these are silent substitutions, additions and deletions, that do not alter the properties and activities of argS.

Further preferred embodiments of the invention are polynucleotides that are at least 70% identical over their entire length to a polynucleotide encoding argS polypeptide having an amino acid sequence set out in Table 1 [SEQ ID NOS: 2 and 4], and polynucleotides that are complementary to such polynucleotides. Alternatively, most highly preferred are polynucleotides that comprise a region that is at least 80% identical over its entire length to a polynucleotide encoding argS polypeptide of the deposited strain and polynucleotides complementary thereto. In this regard, polynucleotides at least 90% identical over their entire length to the same are particularly preferred, and among these particularly preferred polynucleotides, those with at least 95% are especially preferred. Furthermore, those with at least 97% are highly preferred among those with at least 95%, and among these those with at least 98% and at least 99% are particularly highly preferred, with at least 99% being the more preferred.

Preferred embodiments are polynucleotides that encode polypeptides that retain substantially the same biological function or activity as the mature polypeptide encoded by the DNA of Table 1 [SEQ ID NO:1].

The invention further relates to polynucleotides that hybridize to the herein above-described sequences. In this regard, the invention especially relates to polynucleotides that hybridize under stringent conditions to the herein above-described polynucleotides. As herein used, the terms "stringent conditions" and "stringent hybridization conditions" mean hybridization will occur only if there is at least 95% and preferably at least 97% identity between the sequences. An example of stringent hybridization conditions is overnight incubation at 42°C in a solution comprising: 50% formamide, 5x SSC (150mM NaCl, 15mM trisodium citrate), 50 mM sodium phosphate (pH7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 micrograms/ml denatured, sheared salmon sperm DNA, followed by washing the hybridization support in 0.1x SSC at about 65°C. Hybridization and wash conditions are well known and exemplified in Sambrook, *et al.*, Molecular Cloning: A Laboratory Manual, Second Edition, Cold Spring Harbor, N.Y., (1989), particularly Chapter 11 therein.

The invention also provides a polynucleotide consisting essentially of a polynucleotide sequence obtainable by screening an appropriate library containing the complete gene for a polynucleotide sequence set forth in SEQ ID NO: 1 or SEQ ID NO:3 under stringent hybridization conditions with a probe having the sequence of said polynucleotide sequence set forth in SEQ ID NO: or a fragment thereof; and isolating said DNA sequence. Fragments useful for obtaining such a polynucleotide include, for example, probes and primers described elsewhere herein.

As discussed additionally herein regarding polynucleotide assays of the invention, for instance, polynucleotides of the invention as discussed above, may be used as a hybridization probe for RNA, cDNA and genomic DNA to isolate full-length cDNAs and genomic clones encoding argS and to isolate cDNA and genomic clones of other genes that have a high sequence similarity to the argS gene. Such probes generally will comprise at least 15 bases. Preferably, such probes will have at least 30 bases and may have at least 50 bases. Particularly preferred probes will have at least 30 bases and will have 50 bases or less.

For example, the coding region of the argS gene may be isolated by screening using the DNA sequence provided in SEQ ID NO: 1 to synthesize an oligonucleotide probe. A labeled oligonucleotide having a sequence complementary to that of a gene of the invention is then used to screen a library of cDNA, genomic DNA or mRNA to determine which members of the library the probe hybridizes to.

The polynucleotides and polypeptides of the invention may be employed, for example, as research reagents and materials for discovery of treatments of and diagnostics for disease, particularly human disease, as further discussed herein relating to polynucleotide assays.

Polynucleotides of the invention that are oligonucleotides derived from the sequences of SEQ ID NOS: 1 and/or 2 may be used in the processes herein as described, but preferably for PCR, to determine whether or not the polynucleotides identified herein in whole or in part are transcribed in bacteria in infected tissue. It is recognized that such sequences will also have utility in diagnosis of the stage of infection and type of infection the pathogen has attained.

The invention also provides polynucleotides that may encode a polypeptide that is the mature protein plus additional

amino or carboxyl-terminal amino acids, or amino acids interior to the mature polypeptide (when the mature form has more than one polypeptide chain, for instance). Such sequences may play a role in processing of a protein from precursor to a mature form, may allow protein transport, may lengthen or shorten protein half-life or may facilitate manipulation of a protein for assay or production, among other things. As generally is the case *in vivo*, the additional amino acids may be processed away from the mature protein by cellular enzymes.

A precursor protein, having the mature form of the polypeptide fused to one or more prosequences may be an inactive form of the polypeptide. When prosequences are removed such inactive precursors generally are activated. Some or all of the prosequences may be removed before activation. Generally, such precursors are called proproteins.

In sum, a polynucleotide of the invention may encode a mature protein, a mature protein plus a leader sequence (which may be referred to as a preprotein), a precursor of a mature protein having one or more prosequences that are not the leader sequences of a preprotein, or a preproprotein, which is a precursor to a proprotein, having a leader sequence and one or more prosequences, which generally are removed during processing steps that produce active and mature forms of the polypeptide.

## Vectors, host cells, expression

The invention also relates to vectors that comprise a polynucleotide or polynucleotides of the invention, host cells that are genetically engineered with vectors of the invention and the production of polypeptides of the invention by recombinant techniques. Cell-free translation systems can also be employed to produce such proteins using RNAs derived from the DNA constructs of the invention.

For recombinant production, host cells can be genetically engineered to incorporate expression systems or portions thereof or polynucleotides of the invention. Introduction of a polynucleotide into the host cell can be effected by methods described in many standard laboratory manuals, such as Davis et al., *BASIC METHODS IN MOLECULAR BIOLOGY*, (1986) and Sambrook et al., *MOLECULAR CLONING: A LABORATORY MANUAL*, 2nd Ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (1989), such as, calcium phosphate transfection, DEAE-dextran mediated transfection, transvection, microinjection, cationic lipid-mediated transfection, electroporation, transduction, scrape loading, ballistic introduction and infection.

Representative examples of appropriate hosts include bacterial cells, such as streptococci, staphylococci, enterococci *E. coli*, streptomyces and *Bacillus subtilis* cells; fungal cells, such as yeast cells and *Aspergillus* cells; insect cells such as *Drosophila* S2 and *Spodoptera* Sf9 cells; animal cells such as CHO, COS, HeLa, C127, 3T3, BHK, 293 and Bowes melanoma cells; and plant cells.

A great variety of expression systems can be used to produce the polypeptides of the invention. Such vectors include, among others, chromosomal, episomal and virus-derived vectors, *e.g.*, vectors derived from bacterial plasmids, from bacteriophage, from transposons, from yeast episomes, from insertion elements, from yeast chromosomal elements, from viruses such as baculoviruses, papova viruses, such as SV40, vaccinia viruses, adenoviruses, fowl pox viruses, pseudorabies viruses and retroviruses, and vectors derived from combinations thereof, such as those derived from plasmid and bacteriophage genetic elements, such as cosmids and phagemids. The expression system constructs may contain control regions that regulate as well as engender expression. Generally, any system or vector suitable to maintain, propagate or express polynucleotides and/or to express a polypeptide in a host may be used for expression in this regard. The appropriate DNA sequence may be inserted into the expression system by any of a variety of well-known and routine techniques, such as, for example, those set forth in Sambrook *et al.*, *MOLECULAR CLONING, A LABORATORY MANUAL*, (*supra*).

For secretion of the translated protein into the lumen of the endoplasmic reticulum, into the periplasmic space or into the extracellular environment, appropriate secretion signals may be incorporated into the expressed polypeptide. These signals may be endogenous to the polypeptide or they may be heterologous signals.

Polypeptides of the invention can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography, and lectin chromatography. Most preferably, high performance liquid chromatography is employed for purification. Well known techniques for refolding protein may be employed to regenerate active conformation when the polypeptide is denatured during isolation and or purification.

## Diagnostic Assays

This invention is also related to the use of the argS polynucleotides of the invention for use as diagnostic reagents. Detection of argS in a eukaryote, particularly a mammal, and especially a human, will provide a diagnostic method for diagnosis of a disease. Eukaryotes (herein also "individual(s)"), particularly mammals, and especially humans, particularly those infected or suspected to be infected with an organism comprising the argS gene may be detected at the



nucleic acid level by a variety of techniques.

Nucleic acids for diagnosis may be obtained from an infected individual's cells and tissues, such as bone, blood, muscle, cartilage, and skin. Genomic DNA may be used directly for detection or may be amplified enzymatically by using PCR or other amplification technique prior to analysis. RNA or cDNA may also be used in the same ways. Using  
 5 amplification, characterization of the species and strain of prokaryote present in an individual, may be made by an analysis of the genotype of the prokaryote gene. Deletions and insertions can be detected by a change in size of the amplified product in comparison to the genotype of a reference sequence. Point mutations can be identified by hybridizing amplified DNA to labeled argS polynucleotide sequences. Perfectly matched sequences can be distinguished from mismatched duplexes by RNase digestion or by differences in melting temperatures. DNA sequence differences  
 10 may also be detected by alterations in the electrophoretic mobility of the DNA fragments in gels, with or without denaturing agents, or by direct DNA sequencing. See, e.g., Myers et al., *Science*, 230: 1242 (1985). Sequence changes at specific locations also may be revealed by nuclease protection assays, such as RNase and S1 protection or a chemical cleavage method. See, e.g., Cotton et al., *Proc. Natl. Acad. Sci., USA*, 85: 4397-4401 (1985).

Cells carrying mutations or polymorphisms in the gene of the invention may also be detected at the DNA level by a variety of techniques, to allow for serotyping, for example. For example, RT-PCR can be used to detect mutations. It is particularly preferred to use RT-PCR in conjunction with automated detection systems, such as, for example, GeneScan. RNA or cDNA may also be used for the same purpose, PCR or RT-PCR. As an example, PCR primers  
 20 complementary to a nucleic acid encoding argS can be used to identify and analyze mutations. The invention further provides these primers with 1, 2, 3 or 4 nucleotides removed from the 5' and/or the 3' end. These primers may be used for, among other things, amplifying argS DNA isolated from a sample derived from an individual. The primers may be used to amplify the gene isolated from an infected individual such that the gene may then be subject to various techniques for elucidation of the DNA sequence. In this way, mutations in the DNA sequence may be detected and used to diagnose infection and to serotype and/or classify the infectious agent.

The invention further provides a process for diagnosing, disease, preferably bacterial infections, more preferably infections by *Streptococcus pneumoniae*, and most preferably otitis media, conjunctivitis, pneumonia, bacteremia, meningitis, sinusitis, pleural empyema and endocarditis, and most particularly meningitis, such as for example infection of cerebrospinal fluid, comprising determining from a sample derived from an individual a increased level of expression of polynucleotide having the sequence of Table 1 [SEQ ID NO: 1]. Increased or decreased expression of argS poly-  
 30 nucleotide can be measured using any one of the methods well known in the art for the quantitation of polynucleotides, such as, for example, amplification, PCR, RT-PCR, RNase protection, Northern blotting and other hybridization methods.

In addition, a diagnostic assay in accordance with the invention for detecting over-expression of argS protein compared to normal control tissue samples may be used to detect the presence of an infection, for example. Assay  
 35 techniques that can be used to determine levels of a argS protein, in a sample derived from a host are well-known to those of skill in the art. Such assay methods include radioimmunoassays, competitive-binding assays, Western Blot analysis and ELISA assays.

## Antibodies

The polypeptides of the invention or variants thereof, or cells expressing them can be used as an immunogen to produce antibodies immunospecific for such polypeptides. "Antibodies" as used herein includes monoclonal and polyclonal antibodies, chimeric, single chain, simianized antibodies and humanized antibodies, as well as Fab fragments, including the products of an Fab immunoglobulin expression library.

Antibodies generated against the polypeptides of the invention can be obtained by administering the polypeptides or epitope-bearing fragments, analogues or cells to an animal, preferably a nonhuman, using routine protocols. For preparation of monoclonal antibodies, any technique known in the art that provides antibodies produced by continuous cell line cultures can be used. Examples include various techniques, such as those in Kohler, G. and Milstein, C., *Nature* 256: 495-497 (1975); Kozbor et al., *Immunology Today* 4: 72 (1983); Cole et al., pg. 77-96 in *MONOCLONAL ANTIBODIES AND CANCER THERAPY*, Alan R. Liss, Inc. (1985).

Techniques for the production of single chain antibodies (U.S. Patent No. 4,946,778) can be adapted to produce single chain antibodies to polypeptides of this invention. Also, transgenic mice, or other organisms such as other mammals, may be used to express humanized antibodies.

Alternatively phage display technology may be utilized to select antibody genes with binding activities towards the polypeptide either from repertoires of PCR amplified v-genes of lymphocytes from humans screened for possessing  
 55 anti-argS or from naive libraries (McCafferty, J. et al., (1990), *Nature* 348, 552-554; Marks, J. et al., (1992) *Biotechnology* 10, 779-783). The affinity of these antibodies can also be improved by chain shuffling (Clackson, T. et al., (1991) *Nature* 352, 624-628).

If two antigen binding domains are present each domain may be directed against a different epitope - termed 'bispecific' antibodies.

The above-described antibodies may be employed to isolate or to identify clones expressing the polypeptides to purify the polypeptides by affinity chromatography.

Thus, among others, antibodies against argS- polypeptide may be employed to treat infections, particularly bacterial infections and especially otitis media, conjunctivitis, pneumonia, bacteremia, meningitis, sinusitis, pleural empyema and endocarditis, and most particularly meningitis, such as for example infection of cerebrospinal fluid.

Polypeptide variants include antigenically, epitopically or immunologically equivalent variants that form a particular aspect of this invention. The term "antigenically equivalent derivative" as used herein encompasses a polypeptide or its equivalent which will be specifically recognized by certain antibodies which, when raised to the protein or polypeptide according to the invention, interfere with the immediate physical interaction between pathogen and mammalian host. The term "immunologically equivalent derivative" as used herein encompasses a peptide or its equivalent which when used in a suitable formulation to raise antibodies in a vertebrate, the antibodies act to interfere with the immediate physical interaction between pathogen and mammalian host.

The polypeptide, such as an antigenically or immunologically equivalent derivative or a fusion protein thereof is used as an antigen to immunize a mouse or other animal such as a rat or chicken. The fusion protein may provide stability to the polypeptide. The antigen may be associated, for example by conjugation, with an immunogenic carrier protein for example bovine serum albumin (BSA) or keyhole limpet haemocyanin (KLH). Alternatively a multiple antigenic peptide comprising multiple copies of the protein or polypeptide, or an antigenically or immunologically equivalent polypeptide thereof may be sufficiently antigenic to improve immunogenicity so as to obviate the use of a carrier.

Preferably, the antibody or variant thereof is modified to make it less immunogenic in the individual. For example, if the individual is human the antibody may most preferably be "humanized"; where the complementarity determining region(s) of the hybridoma-derived antibody has been transplanted into a human monoclonal antibody, for example as described in Jones, P. et al. (1986), *Nature* 321, 522-525 or Tempest et al., (1991) *Biotechnology* 9, 266-273.

The use of a polynucleotide of the invention in genetic immunization will preferably employ a suitable delivery method such as direct injection of plasmid DNA into muscles (Wolff et al., *Hum Mol Genet* 1992, 1:363, Manthorpe et al., *Hum. Gene Ther.* 1993, 4, 419), delivery of DNA complexed with specific protein carriers (Wu et al., *J Biol Chem.* 1989: 264,16985), coprecipitation of DNA with calcium phosphate (Benvenisty & Reshef, *PNAS USA*, 1986:83,9551), encapsulation of DNA in various forms of liposomes (Kaneda et al., *Science* 1989:243,375), particle bombardment (Tang et al., *Nature* 1992, 356:152, Eisenbraun et al., *DNA Cell Biol* 1993, 12:791) and *in vivo* infection using cloned retroviral vectors (Seeger et al., *PNAS USA* 1984:81,5849).

#### Antagonists and agonists - assays and molecules

Polypeptides of the invention may also be used to assess the binding of small molecule substrates and ligands in, for example, cells, cell-free preparations, chemical libraries, and natural product mixtures. These substrates and ligands may be natural substrates and ligands or may be structural or functional mimetics. See, e.g., Coligan et al., *Current Protocols in Immunology* 1(2): Chapter 5 (1991).

The invention also provides a method of screening compounds to identify those which enhance (agonist) or block (antagonist) the action of argS polypeptides or polynucleotides, particularly those compounds that are bacteriostatic and/or bacteriocidal. The method of screening may involve high-throughput techniques. For example, to screen for agonists or antagonists, a synthetic reaction mix, a cellular compartment, such as a membrane, cell envelope or cell wall, or a preparation of any thereof, comprising argS polypeptide and a labeled substrate or ligand of such polypeptide is incubated in the absence or the presence of a candidate molecule that may be a argS agonist or antagonist. The ability of the candidate molecule to agonize or antagonize the argS polypeptide is reflected in decreased binding of the labeled ligand or decreased production of product from such substrate. Molecules that bind gratuitously, i.e., without inducing the effects of argS polypeptide are most likely to be good antagonists. Molecules that bind well and increase the rate of product production from substrate are agonists. Detection of the rate or level of production of product from substrate may be enhanced by using a reporter system. Reporter systems that may be useful in this regard include but are not limited to colorimetric labeled substrate converted into product, a reporter gene that is responsive to changes in argS polynucleotide or polypeptide activity, and binding assays known in the art.

Another example of an assay for argS antagonists is a competitive assay that combines argS and a potential antagonist with argS-binding molecules, recombinant argS binding molecules, natural substrates or ligands, or substrate or ligand mimetics, under appropriate conditions for a competitive inhibition assay. argS can be labeled, such as by radioactivity or a colorimetric compound, such that the number of argS molecules bound to a binding molecule or converted to product can be determined accurately to assess the effectiveness of the potential antagonist.

Potential antagonists include small organic molecules, peptides, polypeptides and antibodies that bind to a polynucleotide or polypeptide of the invention and thereby inhibit or extinguish its activity. Potential antagonists also may

be small organic molecules, a peptide, a polypeptide such as a closely related protein or antibody that binds the same sites on a binding molecule, such as a binding molecule, without inducing argS-induced activities, thereby preventing the action of argS by excluding argS from binding.

Potential antagonists include a small molecule that binds to and occupies the binding site of the polypeptide thereby preventing binding to cellular binding molecules, such that normal biological activity is prevented. Examples of small molecules include but are not limited to small organic molecules, peptides or peptide-like molecules. Other potential antagonists include antisense molecules (see Okano, *J. Neurochem.* 56: 560 (1991); *OLIGODEOXYNUCLEOTIDES AS ANTISENSE INHIBITORS OF GENE EXPRESSION*, CRC Press, Boca Raton, FL (1988), for a description of these molecules). Preferred potential antagonists include compounds related to and variants of argS.

Each of the DNA sequences provided herein may be used in the discovery and development of antibacterial compounds. The encoded protein, upon expression, can be used as a target for the screening of antibacterial drugs. Additionally, the DNA sequences encoding the amino terminal regions of the encoded protein or Shine-Delgarno or other translation facilitating sequences of the respective mRNA can be used to construct antisense sequences to control the expression of the coding sequence of interest.

The invention also provides the use of the polypeptide, polynucleotide or inhibitor of the invention to interfere with the initial physical interaction between a pathogen and mammalian host responsible for sequelae of infection. In particular the molecules of the invention may be used: in the prevention of adhesion of bacteria, in particular gram positive bacteria, to mammalian extracellular matrix proteins on in-dwelling devices or to extracellular matrix proteins in wounds; to block argS protein-mediated mammalian cell invasion by, for example, initiating phosphorylation of mammalian tyrosine kinases (Rosenshine *et al.*, *Infect. Immun.* 60:2211 (1992); to block bacterial adhesion between mammalian extracellular matrix proteins and bacterial argS proteins that mediate tissue damage and; to block the normal progression of pathogenesis in infections initiated other than by the implantation of in-dwelling devices or by other surgical techniques.

The antagonists and agonists of the invention may be employed, for instance, to inhibit and treat otitis media, conjunctivitis, pneumonia, bacteremia, meningitis, sinusitis, pleural empyema and endocarditis, and most particularly meningitis, such as for example infection of cerebrospinal fluid.

*Helicobacter pylori* (herein *H. pylori*) bacteria infect the stomachs of over one-third of the world's population causing stomach cancer, ulcers, and gastritis (International Agency for Research on Cancer (1994) Schistosomes, Liver Flukes and *Helicobacter Pylori* (International Agency for Research on Cancer, Lyon, France; <http://www.uicc.ch/ecp/ecp2904.htm>). Moreover, the international Agency for Research on Cancer recently recognized a cause-and-effect relationship between *H. pylori* and gastric adenocarcinoma, classifying the bacterium as a Group I (definite) carcinogen. Preferred antimicrobial compounds of the invention (agonists and antagonists of argS) found using screens provided by the invention, particularly broad-spectrum antibiotics, should be useful in the treatment of *H. pylori* infection. Such treatment should decrease the advent of *H. pylori*-induced cancers, such as gastrointestinal carcinoma. Such treatment should also cure gastric ulcers and gastritis.

## Vaccines

Another aspect of the invention relates to a method for inducing an immunological response in an individual, particularly a mammal which comprises inoculating the individual with argS, or a fragment or variant thereof, adequate to produce antibody and/ or T cell immune response to protect said individual from infection, particularly bacterial infection and most particularly *Streptococcus pneumoniae* infection. Also provided are methods whereby such immunological response slows bacterial replication. Yet another aspect of the invention relates to a method of inducing immunological response in an individual which comprises delivering to such individual a nucleic acid vector to direct expression of argS, or a fragment or a variant thereof, for expressing argS, or a fragment or a variant thereof *in vivo* in order to induce an immunological response, such as, to produce antibody and/ or T cell immune response, including, for example, cytokine-producing T cells or cytotoxic T cells, to protect said individual from disease, whether that disease is already established within the individual or not. One way of administering the gene is by accelerating it into the desired cells as a coating on particles or otherwise.

Such nucleic acid vector may comprise DNA, RNA, a modified nucleic acid, or a DNA/RNA hybrid.

A further aspect of the invention relates to an immunological composition which, when introduced into an individual capable or having induced within it an immunological response, induces an immunological response in such individual to a argS or protein coded therefrom, wherein the composition comprises a recombinant argS or protein coded therefrom comprising DNA which codes for and expresses an antigen of said argS or protein coded therefrom. The immunological response may be used therapeutically or prophylactically and may take the form of antibody immunity or cellular immunity such as that arising from CTL or CD4+ T cells.

A argS polypeptide or a fragment thereof may be fused with co-protein which may not by itself produce antibodies, but is capable of stabilizing the first protein and producing a fused protein which will have immunogenic and protective

properties. Thus fused recombinant protein, preferably further comprises an antigenic co-protein, such as lipoprotein D from *Hemophilus influenzae*, Glutathione-S-transferase (GST) or beta-galactosidase, relatively large co-proteins which solubilize the protein and facilitate production and purification thereof. Moreover, the co-protein may act as an adjuvant in the sense of providing a generalized stimulation of the immune system. The co-protein may be attached to either the amino or carboxy terminus of the first protein.

Provided by this invention are compositions, particularly vaccine compositions, and methods comprising the polypeptides or polynucleotides of the invention and immunostimulatory DNA sequences, such as those described in Sato, Y. *et al.* Science 273: 352 (1996).

Also, provided by this invention are methods using the described polynucleotide or particular fragments thereof which have been shown to encode non-variable regions of bacterial cell surface proteins in DNA constructs used in such genetic immunization experiments in animal models of infection with *Streptococcus pneumoniae* will be particularly useful for identifying protein epitopes able to provoke a prophylactic or therapeutic immune response. It is believed that this approach will allow for the subsequent preparation of monoclonal antibodies of particular value from the requisite organ of the animal successfully resisting or clearing infection for the development of prophylactic agents or therapeutic treatments of bacterial infection, particularly *Streptococcus pneumoniae* infection, in mammals, particularly humans.

The polypeptide may be used as an antigen for vaccination of a host to produce specific antibodies which protect against invasion of bacteria, for example by blocking adherence of bacteria to damaged tissue. Examples of tissue damage include wounds in skin or connective tissue caused, e.g., by mechanical, chemical or thermal damage or by implantation of indwelling devices, or wounds in the mucous membranes, such as the mouth, mammary glands, urethra or vagina.

The invention also includes a vaccine formulation which comprises an immunogenic recombinant protein of the invention together with a suitable carrier. Since the protein may be broken down in the stomach, it is preferably administered parenterally, including, for example, administration that is subcutaneous, intramuscular, intravenous, or intradermal. Formulations suitable for parenteral administration include aqueous and non-aqueous sterile injection solutions which may contain anti-oxidants, buffers, bacteriostats and solutes which render the formulation isotonic with the bodily fluid, preferably the blood, of the individual; and aqueous and non-aqueous sterile suspensions which may include suspending agents or thickening agents. The formulations may be presented in unit-dose or multi-dose containers, for example, sealed ampules and vials and may be stored in a freeze-dried condition requiring only the addition of the sterile liquid carrier immediately prior to use. The vaccine formulation may also include adjuvant systems for enhancing the immunogenicity of the formulation, such as oil-in water systems and other systems known in the art. The dosage will depend on the specific activity of the vaccine and can be readily determined by routine experimentation.

While the invention has been described with reference to certain argS protein, it is to be understood that this covers fragments of the naturally occurring protein and similar proteins with additions, deletions or substitutions which do not substantially affect the immunogenic properties of the recombinant protein.

### Compositions, kits and administration

The invention also relates to compositions comprising the polynucleotide or the polypeptides discussed above or their agonists or antagonists. The polypeptides of the invention may be employed in combination with a non-sterile or sterile carrier or carriers for use with cells, tissues or organisms, such as a pharmaceutical carrier suitable for administration to a subject. Such compositions comprise, for instance, a media additive or a therapeutically effective amount of a polypeptide of the invention and a pharmaceutically acceptable carrier or excipient. Such carriers may include, but are not limited to, saline, buffered saline, dextrose, water, glycerol, ethanol and combinations thereof. The formulation should suit the mode of administration. The invention further relates to diagnostic and pharmaceutical packs and kits comprising one or more containers filled with one or more of the ingredients of the aforementioned compositions of the invention.

Polypeptides and other compounds of the invention may be employed alone or in conjunction with other compounds, such as therapeutic compounds.

The pharmaceutical compositions may be administered in any effective, convenient manner including, for instance, administration by topical, oral, anal, vaginal, intravenous, intraperitoneal, intramuscular, subcutaneous, intranasal or intradermal routes among others.

In therapy or as a prophylactic, the active agent may be administered to an individual as an injectable composition, for example as a sterile aqueous dispersion, preferably isotonic.

Alternatively the composition may be formulated for topical application for example in the form of ointments, creams, lotions, eye ointments, eye drops, ear drops, mouthwash, impregnated dressings and sutures and aerosols, and may contain appropriate conventional additives, including, for example, preservatives, solvents to assist drug penetration, and emollients in ointments and creams. Such topical formulations may also contain compatible conventional carriers.

for example cream or ointment bases, and ethanol or oleyl alcohol for lotions. Such carriers may constitute from about 1% to about 98% by weight of the formulation; more usually they will constitute up to about 80% by weight of the formulation.

For administration to mammals, and particularly humans, it is expected that the daily dosage level of the active agent will be from 0.01 mg/kg to 10 mg/kg, typically around 1 mg/kg. The physician in any event will determine the actual dosage which will be most suitable for an individual and will vary with the age, weight and response of the particular individual. The above dosages are exemplary of the average case. There can, of course, be individual instances where higher or lower dosage ranges are merited, and such are within the scope of this invention.

In-dwelling devices include surgical implants, prosthetic devices and catheters, i.e., devices that are introduced to the body of an individual and remain in position for an extended time. Such devices include, for example, artificial joints, heart valves, pacemakers, vascular grafts, vascular catheters, cerebrospinal fluid shunts, urinary catheters, continuous ambulatory peritoneal dialysis (CAPD) catheters.

The composition of the invention may be administered by injection to achieve a systemic effect against relevant bacteria shortly before insertion of an in-dwelling device. Treatment may be continued after surgery during the in-body time of the device. In addition, the composition could also be used to broaden perioperative cover for any surgical technique to prevent bacterial wound infections, especially *Streptococcus pneumoniae* wound infections.

Many orthopaedic surgeons consider that humans with prosthetic joints should be considered for antibiotic prophylaxis before dental treatment that could produce a bacteremia. Late deep infection is a serious complication sometimes leading to loss of the prosthetic joint and is accompanied by significant morbidity and mortality. It may therefore be possible to extend the use of the active agent as a replacement for prophylactic antibiotics in this situation.

In addition to the therapy described above, the compositions of this invention may be used generally as a wound treatment agent to prevent adhesion of bacteria to matrix proteins exposed in wound tissue and for prophylactic use in dental treatment as an alternative to, or in conjunction with, antibiotic prophylaxis.

Alternatively, the composition of the invention may be used to bathe an indwelling device immediately before insertion. The active agent will preferably be present at a concentration of 1µg/ml to 10mg/ml for bathing of wounds or indwelling devices.

A vaccine composition is conveniently in injectable form. Conventional adjuvants may be employed to enhance the immune response. A suitable unit dose for vaccination is 0.5-5 microgram/kg of antigen, and such dose is preferably administered 1-3 times and with an interval of 1-3 weeks. With the indicated dose range, no adverse toxicological effects will be observed with the compounds of the invention which would preclude their administration to suitable individuals.

Each reference disclosed herein is incorporated by reference herein in its entirety. Any patent application to which this application claims priority is also incorporated by reference herein in its entirety.

## EXAMPLES

The examples below are carried out using standard techniques, which are well known and routine to those of skill in the art, except where otherwise described in detail. The examples are illustrative, but do not limit the invention.

### Example 1 Strain selection, Library Production and Sequencing

The polynucleotide having the DNA sequence given in SEQ ID NO: 1 was obtained from a library of clones of chromosomal DNA of *Streptococcus pneumoniae* in *E. coli*. The sequencing data from two or more clones containing overlapping *Streptococcus pneumoniae* DNAs was used to construct the contiguous DNA sequence in SEQ ID NO:1. Libraries may be prepared by routine methods, for example:

Methods 1 and 2 below.

Total cellular DNA is isolated from *Streptococcus pneumoniae* 0100993 according to standard procedures and size-fractionated by either of two methods.

#### Method 1

Total cellular DNA is mechanically sheared by passage through a needle in order to size-fractionate according to standard procedures. DNA fragments of up to 11kbp in size are rendered blunt by treatment with exonuclease and DNA polymerase, and EcoRI linkers added. Fragments are ligated into the vector Lambda ZapII that has been cut with EcoRI, the library packaged by standard procedures and *E. coli* infected with the packaged library. The library is amplified by standard procedures.

**Method 2**

Total cellular DNA is partially hydrolyzed with a one or a combination of restriction enzymes appropriate to generate a series of fragments for cloning into library vectors (e.g., *Rsa*I, *Pa*1I, *Alu*I, *Bsh*1235I), and such fragments are size-fractionated according to standard procedures. *Eco*RI linkers are ligated to the DNA and the fragments then ligated into the vector Lambda ZapII that have been cut with *Eco*RI, the library packaged by standard procedures, and *E. coli* infected with the packaged library. The library is amplified by standard procedures.

**Example 2      argS Characterization**

The enzyme mediated incorporation of radiolabelled amino acid into tRNA may be measured by the aminoacylation method which measures amino acid-tRNA as trichloroacetic acid-precipitable radioactivity from radiolabelled amino acid in the presence of tRNA and ATP (Hughes J, Mellows G and Soughton S, 1980, FEBS Letters, 122:322-324). Thus inhibitors of arginyl tRNA synthetase can be detected by a reduction in the trichloroacetic acid precipitable radioactivity relative to the control. Alternatively the tRNA synthetase catalysed partial PP<sub>i</sub>/ATP exchange reaction which measures the formation of radiolabelled ATP from PP<sub>i</sub> can be used to detect arginyl tRNA synthetase inhibitors (Calender R & Berg P, 1966, Biochemistry, 5, 1681-1690).

## SEQUENCE LISTING

5 (1) GENERAL INFORMATION

(i) APPLICANT: Lawlor, Elizabeth

10 (ii) TITLE OF THE INVENTION: Novel argS

(iii) NUMBER OF SEQUENCES: 4

15 (iv) CORRESPONDENCE ADDRESS:

(A) ADDRESSEE: SmithKline Beecham Corporation

(B) STREET: 709 Swedeland Road

20 (C) CITY: King of Prussia

(D) STATE: PA

(E) COUNTRY: USA

(F) ZIP: 19406-0939

25 (v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Diskette

(B) COMPUTER: IBM Compatible

30 (C) OPERATING SYSTEM: DOS

(D) SOFTWARE: FastSEQ for Windows Version 2.0

(vi) CURRENT APPLICATION DATA:

35 (A) APPLICATION NUMBER:

(B) FILING DATE: 12-SEP-1997

(C) CLASSIFICATION:

40 (vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER: 9619071.5

(B) FILING DATE: 12-SEP-1996

45 (viii) ATTORNEY/AGENT INFORMATION:

(A) NAME: Gimmi, Edward R

(B) REGISTRATION NUMBER: 38,891

(C) REFERENCE/DOCKET NUMBER: P31625

50 (ix) TELECOMMUNICATION INFORMATION:

(A) TELEPHONE: 610-270-4478

(B) TELEFAX: 610-270-5090

55 (C) TELEX:

## (2) INFORMATION FOR SEQ ID NO:1:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1692 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: Genomic DNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

ATGAATACAA	AAGGATTGAT	TGCTAGCGAA	TTGGTTAGCA	TCATTGATAG	CATGGACCAA	60
GAGGTAATTT	TAAAGTTACT	GGAAACCCCT	AAAAACTCAG	AAATGGGGGA	CATCGCTTTC	120
CCTGCTTTTT	CTCTTGCCAA	AGTCGAACGT	AAAGCACCAC	AAATGATTGC	GGCTAAACTG	180
GCTGAAAAAA	TGAACAGCCA	AGCCTTTGAA	AAAGTTGTCT	CAACAGGACC	TTACGTTAAC	240
TTTTTCCTTG	ATAAATCTGC	CATTTCTGCT	CAAGTATTGC	AAGCTGTTAC	CACTGAAAAA	300
GAACACTATG	CTGACCAAAA	TATTGGTAAA	CAAGAAAAATG	TTGTTATCGA	CATGTCTAGT	360
CCGAATATCG	CTAAACCATT	TTTTATTGGC	CACCTGCGTT	CAACTGTTAT	CGGAGATAGC	420
TTGTACACATA	TTTTCCAAAA	AATCGGTTAT	CAAACGGTCA	AGGTCAACCA	TTTGGGAGAC	480
TGGGGTAAAC	AATTTGGGAT	GTTGATTGTT	GCCTACAAAA	AATGGGGCGA	CGAAGAAGCT	540
GTAAAAGCTC	ATCCAATCGA	TGAACTCCTT	AAACTCTATG	TCCGCATCAA	CGCTGAAGCT	600
GAAAATGACC	CTAGCTTGGA	TTANGAAGCG	CGCGAATGGT	TCCGTAAACT	TGAAAATGGA	660
GATGAGGAAG	CTCTCGCTCT	TTGGCAATGG	TTCCGCGATG	AAAGTTTAGT	GGAATTTAAC	720
CGCCTTTACA	ATGAATTGAA	GGTTGAATTT	GACAGCTATA	ACGGAGAAGC	CTTCTACAAT	780
GATAAGATGG	ATGCAGTTGT	AGACATTCTT	TCTGAAAAAG	GACTACTTCT	TGAATCAGAA	840
GGTGCCCAAG	TTGTCAATCT	TGAGAAATAC	GGAATTGAAC	ATCCAGCTCT	CATCAAGAAA	900
TCTGATGGTG	CAACTCTCTA	TATCACACGT	GACTTGGCTG	CAGCCCTTTA	CCGTAAAAAC	960
GAATACGAAT	TTGCTAAATC	TATCTATGTC	GTTGGTCAAG	AACAATCTGC	CCACTTTAAA	1020
CAGCTCAAAG	CTGTCTTGCA	AGAGATGGGC	TACGACTGGA	GTGACGACAT	TACTCACGTT	1080
CCTTTTGGTT	TGGTTACAAA	AGAAGGGAAG	AAACTCTCTA	CTCGTAAAGG	GAATGTCTATC	1140
TTGCTAGAGC	CTACTGTTGC	AGAGGCTGTT	AGCCGTGCCA	AGGTCCAAAT	CGAGGCTAAA	1200
AATCCTGAAC	TAGAAAACAA	AGACCAAGTA	GCACATGCTG	TTGGGGTTGG	AGCCATTAAA	1260
TTCTATGACC	TCAAAAACCGA	CCGTACAAAT	GGATACGACT	TCGACCTAGA	GGCTATGGTA	1320
TCCTTCGAGG	GTGAAACTGG	ACCTTACGTT	CAATATGCCT	ACGCTCGTAT	CCAATCTATC	1380
TTACGCAAAG	CCGATTTCAA	ACCAGAAACA	GCTGGCAACT	ATAGCTTGAA	TGATACTGAA	1440
AGCTGGGAAA	TCATTAAACT	CATTCAAGAC	TTCCACGTA	TTATCAACCG	TGCGGCAGAT	1500
AACTTTGAAC	CTTCTATCAT	TGCTAAATTT	GCAATTAGCC	TAGCTCAATC	CTTTAACAAA	1560
TACTATGCAC	ATACACGTAT	CTTGGATGAA	AGCCCAGAAC	GCGACAGCCG	TCTAGCCCTC	1620
AGCTACGCAA	CCGCAGTCGT	TCTCAAAGAA	GCCCTTCGCT	TGCTTGGAGT	AGAAGCGCCA	1680
GAGAAGATGT	AA					1692

## (2) INFORMATION FOR SEQ ID NO:2:



## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 563 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

```

Met Asn Thr Lys Gly Leu Ile Ala Ser Glu Leu Val Ser Ile Ile Asp
 1             5             10             15
Ser Met Asp Gln Glu Val Ile Leu Lys Leu Leu Glu Thr Pro Lys Asn
20             25             30
Ser Glu Met Gly Asp Ile Ala Phe Pro Ala Phe Ser Leu Ala Lys Val
35             40             45
Glu Arg Lys Ala Pro Gln Met Ile Ala Ala Lys Leu Ala Glu Lys Met
25             50             55             60
Asn Ser Gln Ala Phe Glu Lys Val Val Ala Thr Gly Pro Tyr Val Asn
65             70             75             80
Phe Phe Leu Asp Lys Ser Ala Ile Ser Ala Gln Val Leu Gln Ala Val
30             85             90             95
Thr Thr Glu Lys Glu His Tyr Ala Asp Gln Asn Ile Gly Lys Gln Glu
100             105             110
Asn Val Val Ile Asp Met Ser Ser Pro Asn Ile Ala Lys Pro Phe Phe
35             115             120             125
Ile Gly His Leu Arg Ser Thr Val Ile Gly Asp Ser Leu Ser His Ile
130             135             140
Phe Gln Lys Ile Gly Tyr Gln Thr Val Lys Val Asn His Leu Gly Asp
40             145             150             155             160
Trp Gly Lys Gln Phe Gly Met Leu Ile Val Ala Tyr Lys Lys Trp Gly
165             170             175
Asp Glu Glu Ala Val Lys Ala His Pro Ile Asp Glu Leu Leu Lys Leu
45             180             185             190
Tyr Val Arg Ile Asn Ala Glu Ala Glu Asn Asp Pro Ser Leu Asp Xaa
195             200             205
Glu Ala Arg Glu Trp Phe Arg Lys Leu Glu Asn Gly Asp Glu Glu Ala
50             210             215             220
Leu Ala Leu Trp Gln Trp Phe Arg Asp Glu Ser Leu Val Glu Phe Asn
225             230             235             240
Arg Leu Tyr Asn Glu Leu Lys Val Glu Phe Asp Ser Tyr Asn Gly Glu
55             245             250             255

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50

55

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1462 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: Genomic DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

TTACGTTAAC	TTTTTCCTTG	ATAAATCTGC	CATTTCTGCT	CAAGTATTGC	AAGCTGTTAC	60
CACTGAAAAA	GAACACTATG	CTGACCAAAA	TATTGGTAAA	CAAGAAAATG	TTGTTATCGA	120
CATGTCTAGT	CCGAATATCG	CTAAACCATT	TTTTATTGGC	CACCTGCGTT	CAACTGTTAT	180
CGGAGATAGC	TTGTCACATA	TTTTCCAAAA	AATCGGTTAT	CAAACGGTCA	AGGTCAACCA	240
TTTGGGAGAC	TGGGGTAAAC	AATTGGGAT	GTTGATTGTT	GCCTACAAAA	AATGGGGCGA	300
CGAAGAAGCT	GTAAAAGCTC	ATCCAATCGA	TGAACTCCTT	AAACTCTATG	TCCGCATCAA	360
CGCTGAAGCT	GAAAATGACC	CTAGCTTGGA	TTANGAAGCG	CGCGAATGGT	TCCGTAAACT	420
TGAAAATGGA	GATGAGGAAG	CTCTCGCTCT	TTGGCAATGG	TTCCGCGATG	AAAGTTTAGT	480
GGAATTTAAC	CGCCTTTACA	ATGAATTGAA	GGTTGAATTT	GACAGCTATA	ACGGAGAAGC	540
CTTCTACAAT	GATAAGATGG	ATGCAGTTGT	AGACATTCTT	TCTGAAAAAG	GACTACTTCT	600
TGAATCAGAA	GGTGCCCAAG	TTGTCAATCT	TGAGAAATAC	GGAATTGAAC	ATCCAGCTCT	660
CATCAAGAAA	TCTGATGGTG	CAACTCTCTA	TATCACACGT	GACTTGGCTG	CAGCCCTTTA	720
CCGTAAAAAC	GAATACGAAT	TTGCTAAATC	TATCTATGTC	GTTGGTCAAG	AACAATCTGC	780
CCACTTTAAA	CAGCTCAAAAG	CTGTCTTGCA	AGAGATGGGC	TACGACTGGA	GTGACGACAT	840
TACTCACGTT	CCTTTTGGTT	TGGTTACAAA	AGAAGGGAAG	AAACTCTCTA	CTCGTAAAGG	900
GAATGTCATC	TTGCTAGAGC	CTACTGTTGC	AGAGGCTGTT	AGCCGTGCCA	AGGTCCAAAT	960
CGAGGCTAAA	AATCCTGAAC	TAGAAAAACA	AGACCAAGTA	GCACATGCTG	TTGGGGTTGG	1020
AGCCATTAAA	TTCTATGACC	TCAAAAACCGA	CCGTACAAAT	GGATACGACT	TCGACCTAGA	1080
GGCTATGGTA	TCCTTCGAGG	GTGAAACTGG	ACCTTACGTT	CAATATGCCT	ACGCTCGTAT	1140
CCAATCTATC	TTACGCAAAG	CCGATTTCOA	ACCAGAAACA	GCTGGCAACT	ATAGCTTGAA	1200
TGATACTGAA	AGCTGGGAAA	TCATTAAACT	CATTCAAGAC	TTCCCACGTA	TTATCAACCG	1260
TGCGGCAGAT	AACTTTGAAC	CTTCTATCAT	TGCTAAATTT	GCAATTAGCC	TAGCTCAATC	1320
CTTTAACAAA	TACTATGCAC	ATACACGTAT	CTTGATGAA	AGCCCAGAAC	GCGACAGCCG	1380
TCTAGCCCTC	AGCTACGCAA	CCGCAGTCGT	TCTCAAAGAA	GCCCTTCGCT	TGCTTGGAGT	1440
AGAAGCGCCA	GAGAAGATGT	AA				1462

(2) INFORMATION FOR SEQ ID NO:4:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 486 amino acids

(B) TYPE: amino acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

```

5      Tyr Val Asn Phe Phe Leu Asp Lys Ser Ala Ile Ser Ala Gln Val Leu
      1              5              10              15
      Gln Ala Val Thr Thr Glu Lys Glu His Tyr Ala Asp Gln Asn Ile Gly
      20              25              30
10     Lys Gln Glu Asn Val Val Ile Asp Met Ser Ser Pro Asn Ile Ala Lys
      35              40              45
      Pro Phe Phe Ile Gly His Leu Arg Ser Thr Val Ile Gly Asp Ser Leu
      50              55              60
15     Ser His Ile Phe Gln Lys Ile Gly Tyr Gln Thr Val Lys Val Asn His
      65              70              75              80
      Leu Gly Asp Trp Gly Lys Gln Phe Gly Met Leu Ile Val Ala Tyr Lys
      85              90              95
20     Lys Trp Gly Asp Glu Glu Ala Val Lys Ala His Pro Ile Asp Glu Leu
      100             105             110
      Leu Lys Leu Tyr Val Arg Ile Asn Ala Glu Ala Glu Asn Asp Pro Ser
      115             120             125
25     Leu Asp Xaa Glu Ala Arg Glu Trp Phe Arg Lys Leu Glu Asn Gly Asp
      130             135             140
      Glu Glu Ala Leu Ala Leu Trp Gln Trp Phe Arg Asp Glu Ser Leu Val
      145             150             155             160
30     Glu Phe Asn Arg Leu Tyr Asn Glu Leu Lys Val Glu Phe Asp Ser Tyr
      165             170             175
      Asn Gly Glu Ala Phe Tyr Asn Asp Lys Met Asp Ala Val Val Asp Ile
      180             185             190
35     Leu Ser Glu Lys Gly Leu Leu Leu Glu Ser Glu Gly Ala Gln Val Val
      195             200             205
      Asn Leu Glu Lys Tyr Gly Ile Glu His Pro Ala Leu Ile Lys Lys Ser
      210             215             220
40     Asp Gly Ala Thr Leu Tyr Ile Thr Arg Asp Leu Ala Ala Ala Leu Tyr
      225             230             235             240
      Arg Lys Asn Glu Tyr Glu Phe Ala Lys Ser Ile Tyr Val Val Gly Gln
      245             250             255
45     Glu Gln Ser Ala His Phe Lys Gln Leu Lys Ala Val Leu Gln Glu Met
      260             265             270
      Gly Tyr Asp Trp Ser Asp Asp Ile Thr His Val Pro Phe Gly Leu Val
      275             280             285
50     Thr Lys Glu Gly Lys Lys Leu Ser Thr Arg Lys Gly Asn Val Ile Leu
      290             295             300
      Leu Glu Pro Thr Val Ala Glu Ala Val Ser Arg Ala Lys Val Gln Ile
      305             310             315             320
55

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Glu Ala Lys Asn Pro Glu Leu Glu Asn Lys Asp Gln Val Ala His Ala  
 325 330 335  
 5 Val Gly Val Gly Ala Ile Lys Phe Tyr Asp Leu Lys Thr Asp Arg Thr  
 340 345 350  
 Asn Gly Tyr Asp Phe Asp Leu Glu Ala Met Val Ser Phe Glu Gly Glu  
 355 360 365  
 10 Thr Gly Pro Tyr Val Gln Tyr Ala Tyr Ala Arg Ile Gln Ser Ile Leu  
 370 375 380  
 Arg Lys Ala Asp Phe Lys Pro Glu Thr Ala Gly Asn Tyr Ser Leu Asn  
 385 390 395 400  
 15 Asp Thr Glu Ser Trp Glu Ile Ile Lys Leu Ile Gln Asp Phe Pro Arg  
 405 410 415  
 Ile Ile Asn Arg Ala Ala Asp Asn Phe Glu Pro Ser Ile Ile Ala Lys  
 420 425 430  
 20 Phe Ala Ile Ser Leu Ala Gln Ser Phe Asn Lys Tyr Tyr Ala His Thr  
 435 440 445  
 Arg Ile Leu Asp Glu Ser Pro Glu Arg Asp Ser Arg Leu Ala Leu Ser  
 450 455 460  
 25 Tyr Ala Thr Ala Val Val Leu Lys Glu Ala Leu Arg Leu Leu Gly Val  
 465 470 475 480  
 Glu Ala Pro Glu Lys Met  
 485  
 30

### 35 Claims

1. An isolated polynucleotide comprising a polynucleotide sequence selected from the group consisting of:
  - (a) a polynucleotide having at least a 70% identity to a polynucleotide encoding a polypeptide comprising the amino acid sequence of SEQ ID NO:2;
  - (b) a polynucleotide having at least a 70% identity to a polynucleotide encoding the same mature polypeptide expressed by the argS gene contained in the *Streptococcus pneumoniae* of the deposited strain;
  - (c) a polynucleotide encoding a polypeptide comprising an amino acid sequence which is at least 70% identical to the amino acid sequence of SEQ ID NO:2;
  - (d) a polynucleotide which is complementary to the polynucleotide of (a), (b) or (c); and
  - (e) a polynucleotide comprising at least 15 sequential bases of the polynucleotide of (a), (b), (c) or (d).
2. The polynucleotide of Claim 1 wherein the polynucleotide is DNA.
3. The polynucleotide of Claim 1 wherein the polynucleotide is RNA.
4. The polynucleotide of Claim 2 comprising the nucleic acid sequence set forth in SEQ ID NO:1.
5. The polynucleotide of Claim 2 comprising nucleotide 1 to 1689 set forth in SEQ ID NO:1.
6. The polynucleotide of Claim 2 which encodes a polypeptide comprising the amino acid sequence of SEQ ID NO:2.
7. A vector comprising the polynucleotide of Claim 1.

8. A host cell comprising the vector of Claim 7.
9. A process for producing a polypeptide comprising: expressing from the host cell of Claim 8 a polypeptide encoded by said DNA.
10. A process for producing a argS polypeptide or fragment comprising culturing a host of claim 8 under conditions sufficient for the production of said polypeptide or fragment.
11. A polypeptide comprising an amino acid sequence which is at least 70% identical to the amino acid sequence of SEQ ID NO:2.
12. A polypeptide comprising an amino acid sequence as set forth in SEQ ID NO:2.
13. An antibody against the polypeptide of claim 11.
14. An antagonist which inhibits the activity or expression of the polypeptide of claim 11.
15. A method for the treatment of an individual in need of argS polypeptide comprising: administering to the individual a therapeutically effective amount of the polypeptide of claim 11.
16. A method for the treatment of an individual having need to inhibit argS polypeptide comprising: administering to the individual a therapeutically effective amount of the antagonist of Claim 14.
17. A process for diagnosing a disease related to expression or activity of the polypeptide of claim 11 in an individual comprising:
- (a) determining a nucleic acid sequence encoding said polypeptide, and/or
  - (b) analyzing for the presence or amount of said polypeptide in a sample derived from the individual.
18. A method for identifying compounds which interact with and inhibit or activate an activity of the polypeptide of claim 11 comprising:
- contacting a composition comprising the polypeptide with the compound to be screened under conditions to permit interaction between the compound and the polypeptide to assess the interaction of a compound, such interaction being associated with a second component capable of providing a detectable signal in response to the interaction of the polypeptide with the compound;
  - and determining whether the compound interacts with and activates or inhibits an activity of the polypeptide by detecting the presence or absence of a signal generated from the interaction of the compound with the polypeptide.
19. A method for inducing an immunological response in a mammal which comprises inoculating the mammal with argS polypeptide of claim 11, or a fragment or variant thereof, adequate to produce antibody and/or T cell immune response to protect said animal from disease.
20. A method of inducing immunological response in a mammal which comprises delivering a nucleic acid vector to direct expression of argS polypeptide of claim 11, or fragment or a variant thereof, for expressing said argS polypeptide, or a fragment or a variant thereof *in vivo* in order to induce an immunological response to produce antibody and/ or T cell immune response to protect said animal from disease.

(19)



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(54) **Arginyl tRNA synthase**

(57) The invention provides Arginyl tRNA synthetase (argS) polypeptides and DNA (RNA) encoding argS polypeptides and methods for producing such

polypeptides by recombinant techniques. Also provided are methods for utilizing argS polypeptides to screen for antibacterial compounds.

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# PARTIAL EUROPEAN SEARCH REPORT

Application Number

which under Rule 45 of the European Patent Convention EP 97 3C 7006 shall be considered, for the purposes of subsequent proceedings, as the European search report

DOCUMENTS CONSIDERED TO BE RELEVANT			
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X	WO 96 08582 A (BERGERON, M.G. & OULETTE, M.) 21 March 1996 (1996-03-21) SEQ ID NO: 31 * page 88 - page 89 * * claims 1,2,4,52,53,55,118,120,121,123-125 * ---	1-3, 7-10,17	C12N15/52 C12N9/00 C07K16/40 G01N33/50 G01N33/53 C12Q1/68 A61K38/53 A61K39/09
Y	VON DER HAAR, F. ET AL.: "Target Directed Drug Synthesis as Possible Targets" ANGEWANDTE CHEMIE, vol. 20, no. 3, March 1981 (1981-03), pages 217-223, XP002112772 * the whole document *	1-13,15, 17-20	
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INCOMPLETE SEARCH			
<p>The Search Division considers that the present application, or one or more of its claims, does/do not comply with the EPC to such an extent that a meaningful search into the state of the art cannot be carried out, or can only be carried out partially, for these claims.</p> <p>Claims searched completely :</p> <p>Claims searched incompletely :</p> <p>Claims not searched :</p> <p>Reason for the limitation of the search:</p> <p>see sheet C</p>			
Place of search		Date of completion of the search	Examiner
BERLIN		27 August 1999	Fuchs, U
CATEGORY OF CITED DOCUMENTS			
<p>X : particularly relevant if taken alone</p> <p>Y : particularly relevant if combined with another document of the same category</p> <p>A : technological background</p> <p>O : non-written disclosure</p> <p>P : intermediate document</p> <p>T : theory or principle underlying the invention</p> <p>E : earlier patent document, but published on, or after the filing date</p> <p>D : document cited in the application</p> <p>L : document cited for other reasons</p> <p>&amp; : member of the same patent family, corresponding document</p>			

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INCOMPLETE SEARCH  
SHEET C

Application Number

EP 97 30 7006

Although claims 15, 19 and 20 are directed to a method of treatment of the human/animal body (Article 52(4) EPC), the search has been carried out and based on the alleged effects of the compound/composition.

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Claim(s) not searched:  
14, 16

Reason for the limitation of the search:

Claims 14 and 16, pertaining to an antagonist, could not be searched as the subject-matter has not been sufficiently disclosed.



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## PARTIAL EUROPEAN SEARCH REPORT

Application Number  
EP 97 30 7006

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**ANNEX TO THE EUROPEAN SEARCH REPORT  
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The members are as contained in the European Patent Office EDP file on  
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27-08-1999

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